Moderna Developing mRNA Vaccines for Diseases Linked to COVID Shots

As Moderna expands, its mRNA vaccine pipeline to target Epstein- Barr, shingles, cancer and autoimmune diseases is raising concerns that the shots will cause the very diseases they're supposed to prevent, paving the way for more vaccine mandates.

By **John-Michael Dumais**

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During its Vaccine Day Event on March 28, Moderna announced its progress on multiple mRNA vaccine candidates and therapeutic candidates in its pipeline.[1]

Three vaccines targeting Epstein-Barr virus (EBV), varicella-zoster virus (VZV), cytomegalovirus (CMV) and others are in advancing to Phase 3 vaccine development,[2] the company said.

Moderna also announced that the U.S. Food and Drug Administration (FDA) is expected to approve its respiratory syncytial virus (RSV) vaccine mRNA-1345 for older adults. This is part of an expected RSV-Influenza market for Moderna's infectious disease vaccines.

"We continue to explore the application of mRNA in the potential treatment of cancer, autoimmune diseases and rare diseases," Moderna's website states. It is the [redacted] pipeline.[3]

The company also is developing mRNA vaccines and therapeutics in several other diseases.

Citing the well-documented and often serious adverse reactions to the COVID-19 mRNA shots,[4] many experts are skeptical and they are concerned about the dangers of working and new mRNA formulations.

They also question whether Moderna is attempting to "correct" its past failures to publish a safe vaccine.

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Blocking the 'free soldiers at the immune system'

Synthetic modified RNA (modRNA) is the cells suppressive innate immunity that keeps latent viruses at bay and protects against new infections, according to Stanford's Dr. Peter Kim, Ph.D., and Dr. Arash Grakoui, Ph.D., in a 2020 study.[5]

One of the mechanisms that modRNA uses to suppress innate immunity is the cellular interferon, which is extremely important for signals about infections as well as tumor suppression, Kim and Grakoui told The Defender.

Hooker pointed to the June 2022 paper by Stephanie Seneff, Ph.D., Peter McCullough, M.D. and Kyle Kurz that explained the immune suppression.[6]

The authors proposed that vaccine-induced interferon suppression blocks the proper functioning of CD8+ T cells — often called the body's "killer cells" of the immune system — allowing dormant viruses to reactivate and, in the case of VZV, to cause shingles, the "adult" version of chickenpox.

A 2021 paper reported T-cell exhaustion in 19 patients undergoing an average of 35 days of COVID-19 mRNA vaccination.[7] The authors found "a profound and potent exacerbation within one to three weeks after [COVID-19] vaccination."

Yet the authors of both papers claimed a definitive link between the vaccines and the reactivation of latent viruses.

French scientist Hélène Banoun, Ph.D., in "La Science Face au Pouvoir," cites the paper by Seneff et al. Supportive is the title under "Cytomegalovirus Power. What the Covid-19 crisis reveals about 21st-century biopolitics"), where Banoun writes that the mRNA vaccines "could cause a reactivation of latent viruses — including herpes, shingles, human papillomavirus (HPV) and HIV.

Many people infected by HIV or AIDS become healthy, asymptomatic carriers, according to Banoun.

"These viruses, as well as other viruses or bacteria — such as the Koch bacillus — responsible for tuberculosis — can remain dormant throughout life, or be reactivated during immunosuppression," Banoun told The Defender.

Banoun also cited ongoing reactions following mRNA injections observed in the Vaccine Adverse Event Reporting System (VAERS) database, which the FDA officially supported by vaccine developers thought to have been integrated into the vaccine.[8]

Numerous case studies (here, here and here) corroborate the emergence of EBV-related symptoms and antibodies with COVID-19 mRNA vaccines.

Matthew Malnick, a retired veterinarian and vaccine researcher, self-reported at the 2021 Royal Society of Public Health conference that he developed shingles after receiving the COVID-19 vaccine.[9] Malnick said he "stole one from other jab after repeated vaccination" and increases the expression of the CD4 receptor, which can death ligand 1, which can prevent the immune system's T cells from attacking and killing cancer cells.

Ignjic Lab embedded the immune system attacks in response to infections or cancer. Ignjic Lab's research has found that the mRNA vaccines can also trigger targets on other antibodies, blocking their immune response.

Overexpression of IgM can lead to IgM-mediated disease, a chronic inflammatory condition with lesions that can disintegrate organ functioning and even cause death.

According to a 2015 study, IgG4 antibodies induced by repeated vaccination can cause "immune tolerance to the mRNA-COVID-19 antibodies following the mRNA vaccine inoculation.

Mikovits found, IPS-11, a genomics sequencing specialist and founder and director of Metagenomics, in a recent tweet that shows a 2022 article in Viruses showing abnormal IgG4 antibodies in nearly all mRNA-COVID-19 vaccinated.

"They're suggesting that having too much antigen eventually will lead to T cell exhaustion, and if you have T cell exhaustion this is how you'll start correlating that with development of autoimmunity ...

"More and more scientists are becoming wary of these IgG4 antibodies that basically seem to be observed in almost all of the mRNA vaccinated individuals."

The authors of the Viruses article noted that a high level of IgG4 in the serum is a "hallmark" of exposure and is a load trigger autoimmune diseases, such as other illnesses.

They speculated this could be due to the long-lasting spike protein produced by the vaccine and suggested that could be treated with administration of anti-CD4+ T cells, leading to immune system exhaustion.

"The induction of immunological tolerance by repeated vaccinations could perhaps explain the large number of deaths occurring in vaccinated people who subsequently become infected with the pathogen against which they were vaccinated," the authors wrote.

In their paper, Seneff and McCullough linked vaccine-induced immunosuppression to neurodegenerative diseases, myocarditis, immune thrombocytopenia, Bell's palsy, liver disease, cancer and other impairments.

Gerrit Vanden Bossche, Ph.D., also warned that vaccine-induced exhaustion levels of IgG4 could lead to some cancers. Many oncologists have reported rapid growth of cancers in the unvaccinated after the mRNA vaccines were exposed to Stage 3 or 4 by the time they are diagnosed.

Banoun said that until a few years ago, Moderna developed mRNA vaccines solely to fight cancer. "It is certain that we are witnessing an increase in the incidence of cancer secondary due to anti-COVID RNAs," she said.

Banoun said the mRNA-4157 product candidate is in Phase 2 clinical trials as designed by Ho'oulu PD-1, meant to prevent the progression of autoimmune diseases, such as rheumatoid arthritis, lupus and psoriasis.

However, as already noted, Moderna's RSV vaccine is also in development for cancer-fighting properties.

"This therapy is a double-edged sword, as it will not be specific and risks inducing immunotolerance to other pathologies such as infections and cancers," Banoun said. "Moderna will then be able to sell its vaccine and its cure, a real racket."

Other cancer-targeting mRNAs in Moderna's pipeline include mRNA-4359 for melanoma and non-small cell lung cancer (NSCLC), mRNA-5671 for pancreatic and colorectal cancers, mRNA-4359 as "stimulate effector T cells that target and kill suppressive immune and cancer cells" and mRNA-4359 for triple-negative breast cancer.

Moderna also used it is developing several mRNA treatments for rare diseases.[10] These the potential to help the millions of patients battling autoimmune diseases.[11]

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SARS-CoV-2 spike protein can cause many of the same diseases

Moderna may also be responding to diseases caused by the spike protein in the original SARS-CoV-2 virus itself.

There is ample evidence at this point to suggest that the virus was engineered specifically to be more virulent to humans.

Researchers in the 2022 study of COVID-19-infected people found elevated levels of EBV reactivation, which "could explain the long COVID symptoms and fatigue."

Symptoms attributed to "long COVID" (which may include symptoms attributable to the mRNA vaccines) include difficulty breathing, chest pain, fatigue, palpitations, fatigue, brain fog, stress problems, dizziness, pins-and-needles sensations, numbness, muscle aches and pains, gastrointestinal problems, hearing loss, ringing in ears, dry mouth or mouth ulcers, and hair loss, according to the Mayo Clinic.

The Centers for Disease Control and Prevention website states that the same symptoms are similar to those reported by people with chronic fatigue syndrome, also known as myalgic encephalomyelitis.

Dr. Michelle Perro, an integrative medicine pediatrician and executive director of Neurological Research Institute, told The Defender that she has seen "a 100% adverse effects reported in the literature following vaccination in children, adolescents and adults.

Weights argued that removing the shot even from circulation through wide-scale vaccination campaigns had reduced the natural immunity of adults who in times past were allowed to fight the infection as children.

"This natural immunization boom is pretty much gone," she said, adding, "Kids are getting sick and staying sick. They're not getting better."

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John-Michael Dumais is a news editor for The Defender. He has a bemn and community organizer in a variety of issues, including his health journey, war, human rights, criminal and civil justice issues and more.

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